FORM PTO-1449 U.S. Department of Commerce Patent and Trademark Office List of Documents Cited by Applicant	Attorney Docket No.: 1386/19	Serial No.: 10/806,899
	Applicant(s): Petrou et al.	
	Filing Date: March 23, 2004	Group:

U.S. PATENT DOCUMENTS

Examiner Initial	No.	Document Number	Date	Name	Class	Subclass	Filing date if Appropriate

FOREIGN PATENT DOCUMENTS

		Document Number	Date	Country	Name of Patentee or Applicant	Translation Yes | No
STK	1.	WO 02/50096	6/27/2002	PCT	Bionomics Limited	

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

STK	2.	Annegers, "The Epidemiology of Epilepsy," The Treatment of Epilepsy: Principles and Practice, **2nd ed**: 165-172 (1996).
	3.	Berkovic et al., "Concepts of absence epilepsies: Discrete syndromes or biological continuum?" Neurology, **37(6)**: 993-1000 (June 1987).
	4.	Berkovic et al., "The epilepsies: specific syndromes or a neurobiological continuum?" Epileptic Seizures and Syndromes, pp. 25-37 (1994).
	5.	Bourgeois, "Chronic Management of Seizures in the Syndromes of Idiopathic Generalized Epilepsy," Epilepsia, **44(Suppl. 2)**:27-32 (2003).
	6.	Claes et al., "De Novo Mutations in the Sodium-Channel Gene SCN1A Cause Severe Myoclonic Epilepsy of Infancy," American Journal of Human Genetics, **68**:1327-1332 (2001).
	7.	Commission on Classification and Terminology of the International League Against Epilepsy, "Proposal for Revised Classification of Epilepsies and Epileptic Syndromes," Epilepsia, **30(4)**: 389-399 (1989).
	8.	Escayg et al., "Mutations of SCN1A, Encoding a Neuronal Sodium Channel, in two Families with GEFS+2," Nature Genetics, **24**: 343-345 (April 2000).

FORM PTO-1449 U.S. Department of Commerce Patent and Trademark Office List of Documents Cited by Applicant	Attorney Docket No.: 1386/19	Serial No.: 10/806,899
	Applicant(s): Petrou et al.	
	Filing Date: March 23, 2004	Group:

STK	9.	Gardiner, "Impact of our Understanding of the Genetic Aetiology of Epilepsy," Journal of Neurology, **247**: 327-334 (2000).
	10.	Mulley et al., "Channelopathies as a Genetic Cause of Epilepsy," Current Opinion in Neurology, **16**:171-176 (2003).
	11.	Nabbout et al., "Spectrum of SCN1A Mutations in Severe Myoclonic Epilepsy of Infancy," Neurology, **60**:1961-1967 (June 2003).
	12.	Ohmori et al., "Significant Correlation of the SCN1A Mutations and Severe Myoclonic Epilepsy in Infancy," Biochemical and Biophysical Research Communications, **295**:17-23 (2002).
	13.	Reutens et al., "Idiopathic Generalized Epilepsy of Adolescence: Are the Syndromes Clinically Distinct?" Neurology, **45**:1469-1476 (August 1995).
	14.	Scheffer et al., "Generalized Epilepsy with Febrile Seizures Plus: A Genetic Disorder with Heterogeneous Clinical Phenotypes," Brain, **120**:479-490 (1997).
	15.	Scheffer et al., "The Genetics of Human Epilepsy," TRENDS in Pharmacological Science, **24(8)**: 428-433 (August 2003).
	16.	Singh et al., "Generalized Epilepsy with Febrile Seizures Plus: A Common Childhood-Onset Genetic Epilepsy Syndrome," Annals of Neurology, **45(1)**: 75-81 (1999).
	17.	Singh et al., "Severe Myoclonic Epilepsy of Infancy: Extended Spectrum of GEFS?" Epilepsia, **42(7)**: 837-844 (2001).
	18.	Sugawara et al., "Frequent Mutations of SCN1A in Severe Myoclonic Epilepsy in Infancy," Neurology, **58**: 1122-1124 (2002).
↓	19.	Veggiotti et al., "Generalized Epilepsy with Febrile Seizures plus and Severe Myoclonic Epilepsy in Infancy: a case report of two Italian families," Epileptic Discord, **3**: 29-32 (2001).

EXAMINER /Stephen Kapushoc/ DATE CONSIDERED 11/13/2006

*Examiner Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

FORM PTO-1449 U.S. Department of Commerce Patent and Trademark Office List of Documents Cited by Applicant	Attorney Docket No.: 1386/19	Serial No.: 10/806,899
	Applicant(s): Petrou et al.	
	Filing Date: March 23, 2004	Group:

U.S. PATENT DOCUMENTS

Examiner Initial	No.	Document Number	Date	Name	Class	Subclass	Filing date if Appropriate

FOREIGN PATENT DOCUMENTS

		Document Number	Date	Country	Name of Patentee or Applicant	Translation Yes | No

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

STK	1.	International Search Report for PCT/AU2004/000295 dated May 14, 2004.
	2.	Examiner's First Report for Australian Patent Application No. 2004200978 dated May 6, 2004.
	3.	Cannon, *Sodium Channel Gating: No Margin for Error*, Neuron **34**:853-858 (June 13, 2002).
	4.	Fujiwara et al., *Mutations of sodium channel α subunit type 1 (SCN1A) in intractable childhood epilepsies with frequent generalized tonic-clonic seizures*, Brain **126**:531-546 (2003).
	5.	Lerche et al., *Ion Channels and Epilepsy*, Am. J. of Med. Genetics **106**:146-159 (2001).
	6.	Madia et al., *No evidence of GABRG2 mutations in severe myoclonic epilepsy of infancy*, Epilepsy Research **53**:196-200 (2003).
	7.	Malacarne et al., *Lack of SCN1A Mutations in Familial Febrile Seizures*, Epelepsia **43(5)**:559-562 (2002).
↓	8.	Ohmori, et al., *Significant correlation of the SCN1A mutations and severe myoclonic epilepsy in infancy*, Biochemical and Biophysical Research Communications **295**:17-23 (2002).

FORM PTO-1449 U.S. Department of Commerce Patent and Trademark Office List of Documents Cited by Applicant	Attorney Docket No.: 1386/19	Serial No.: 10/806,899
	Applicant(s): Petrou et al.	
	Filing Date: March 23, 2004	Group:

STK	9.	Spampanato et al., *Generalized Epilepsy with Febrile Seizures Plus Type 2 Mutation W1204R Alters voltage-Dependent Gating of $Na_v1.1$ Sodium Channels*, Neuroscience **116**:37-48 (2003).
STK	10.	Sugawara et al., *Frequent mutations of SCN1A in severe myoclonic epilepsy in infancy*, Neurology **58**:1122-1124 (2002).

EXAMINER /Stephen Kapushoc/ DATE CONSIDERED 11/13/2006

*Examiner Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

FORM PTO-1449 U.S. Department of Commerce Patent and Trademark Office List of Documents Cited by Applicant	Attorney Docket No. 1386/19	Serial No. 10/806,899
	Applicant(s): Petrou et al.	
	Filing Date: March 23, 2004	Group: 1614

U.S. PATENT DOCUMENTS

Examiner Initial		Document Number	Date	Name	Class	Subclass	Filing date if Appropriate

FOREIGN PATENT DOCUMENTS

		Document Number	Date	Country	Name of Patentee or Applicant	Translation Yes | No
STK	A	WO01/38564 A2	05/31/2001	WIPO	McGill University, Canada	No

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

EXAMINER /Stephen Kapushoc/ DATE CONSIDERED 11/13/2006

*Examiner Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.